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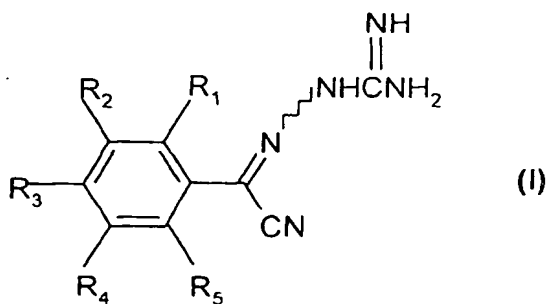
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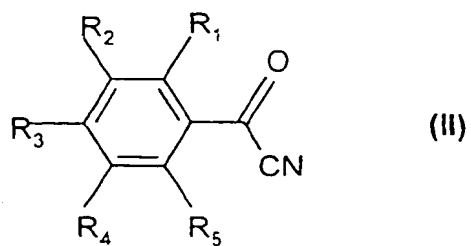
(54) Process for preparing substituted Benzoyl Cyanide Amidinohydrazones

(57) The present invention provides a process for preparing a compound of the general formula (I):



wherein R_1 to R_5 are independently selected from hydrogen, halogen, C_{1-6} alkyl, C_{2-6} alkenyl, C_{2-6} alkynyl or C_{1-6} alkoxy, all optionally substituted by one or more of halogen, hydroxy and aryl groups, as well as being independently selected from amino, mono- or disubstituted amino, alkenyloxy, acyl, acyloxy, cyano, nitro, aryl and alkylthio groups, which process comprises reacting a compound of the general formula (II):

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wherein R₁ to R₅ are as defined in formula (I), with aminoguanidine bicarbonate in a mixture of a water-soluble solvent and polyphosphoric acid.

Description

Field of the Invention:

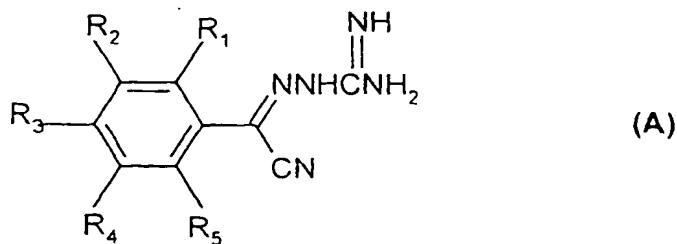
[0001] The present invention relates to a method for preparing substituted benzoyl cyanide amidinohydrazones. More particularly, the present invention pertains to the preparation of these compounds by a reaction of substituted benzoyl cyanides with aminoguanidine in the presence of polyphosphoric acid.

Background of the Invention:

[0002] It is known that certain 3,5-diamino-6-(substituted phenyl)-1,2,4-triazines are active in the treatment of CNS disorders, such as psychiatric and neurological disorders, and are also useful as anticonvulsants, for example in the treatment of epilepsy. These triazines are also non-depressant at therapeutic dose levels and are therefore advantageous as compared with depressant anti-epileptics such as phenobarbitone.

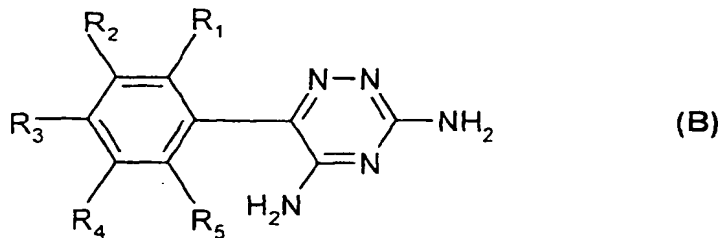
[0003] A particularly preferred compound of this type is 3,5-diamino-6-(2,3-dichlorophenyl)-1,2,4-triazine (e.g. European Patent 21121).

[0004] A known process for preparing 3,5-diamino-6-(substituted phenyl)-1,2,4-triazines, comprises reacting substituted benzoyl cyanides with aminoguanidine in aqueous solutions of strong acids such as nitric acid (J.A. Settepani and A.B. Borkovec, J. Heterocycl. Chem., 1966, 3, 188, U.S. Patent Nos. 3,637,688; 4,486,354; 4,602,017; 4,649,139, R.W.A. Rees et al, J. Med. Chem., 1972, 15, 859; European Patent Nos. 21,121; 142,306; 247,892, Israeli Patent Nos. 60,201; 73,332; 82,710) and sulfuric acid (European Patent No. 247,892, Israeli Patent No. 82,710) to produce amidinohydrazone of the general formula (A):



wherein R_1 is halogen, C_{1-4} alkyl or trifluoromethyl, R_2 is hydrogen, halogen, C_{1-4} alkyl or trifluoromethyl, or R_1 and R_2 form a $-CH=CH-CH=CH-$ group optionally substituted by a halogen atom or a C_{1-4} alkyl or trifluoromethyl group, R_3 and R_4 are independently selected from hydrogen, halogen, C_{1-4} alkyl or trifluoromethyl groups and R_5 is hydrogen, methyl or fluorine.

[0005] 3,5-Diamino-6-(substituted phenyl)-1,2,4-triazines (compounds of the general formula (B):



wherein R_1 - R_5 are as hereinbefore defined) are obtained by ring closure of a compound of the formula A.

[0006] The ring closure is normally carried out by refluxing in an alcohol, such as methanol, ethanol, 1-propanol and the like, in the presence of a strong base, such as sodium hydroxide, potassium hydroxide and the like.

[0007] However, the reaction of substituted benzoyl cyanides with aminoguanidine in the aqueous solutions of strong acids is very slow and usually from 2 to 21 days are required to complete this reaction. On the other hand, substituted benzoyl cyanides are hydrolyzed at the reaction conditions, and overall yield of the triazines is low (from 2 to 41%) (Table).

Table

Compounds of the formula (A)	Solvent of the reaction	Temperature of the reaction	Time of the reaction	Yield of the compound of the formula (B)
R ⁶ =R ⁷ =Cl R ⁸ =R ⁹ =R ¹⁰ =H	a) DMSO and 8N aqueous nitric acid	25°C	7 days	15.6% *
	b) Acetonitrile and 63% aqueous sulfuric acid	20-30°C	48 hours	41% **
R ⁶ =R ⁷ =R ⁹ =Cl R ⁸ =R ¹⁰ =H	DMSO and 8N aqueous nitric acid	20-30°C	21 days	1.6% *
R ⁶ =R ⁸ =Cl R ⁷ =R ⁹ =R ¹⁰ =H	DMSO and 8N aqueous nitric acid	25°C	24 hours	8% ***
R ⁷ =R ⁸ =Cl R ⁶ =R ⁹ =R ¹⁰ =H	DMSO and 8N aqueous nitric acid	25°C	24 hours	23% ***
R ⁸ =Cl R ⁶ =R ⁷ =R ⁹ =R ¹⁰ =H	DMSO and 8N aqueous nitric acid	25°C	24 hours	37% ***

* U.S. Patent No. 4,486,354

** European Patent No. 247,892

*** R.W.A. Rees et al, *J. Med. Chem.*, 1972, 15, 859.**Summary of the Invention:**

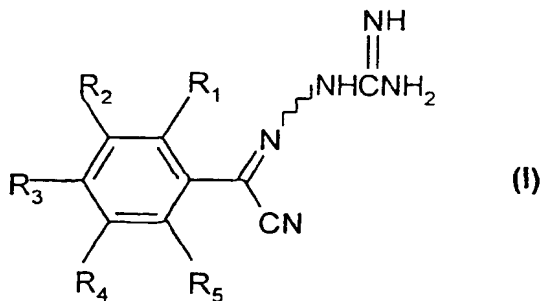
[0008] As a result of research to solve the drawbacks of the prior art methods, the present inventors have found the method for preparing the substituted benzoyl cyanide amidinohydrazones by a reaction of substituted benzoyl cyanides with aminoguanidine in a mixture of polyphosphoric acid and a solvent in the absence of water which minimizes the hydrolysis of the benzoyl cyanides.

[0009] According to the present invention substituted benzoyl cyanide amidinohydrazones can be prepared in higher yield after the reaction of substituted benzoyl cyanide with aminoguanidine in a mixture of polyphosphoric acid and acetonitrile for 24 hours.

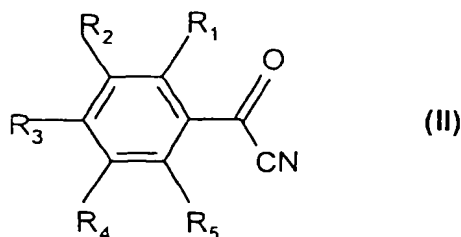
[0010] Other objects and advantages of the present invention will be apparent from the following description.

Detailed Description of the Invention:

[0011] More specifically, the present invention provides a process for preparing a compound of the general formula (I):



wherein R₁ to R₅ are independently selected from hydrogen, halogen, C₁₋₆ alkyl, C₂₋₆ alkenyl, C₂₋₆ alkynyl or C₁₋₆ alkoxy, all optionally substituted by one or more of halogen, hydroxy and aryl groups, as well as being independently selected from amino, mono- or disubstituted amino, alkenyloxy, acyl, acyloxy, cyano, nitro, aryl and alkylthio groups, which process comprises reacting a compound of the general formula (II):



wherein R_1 to R_5 are as defined in formula (I), with aminoguanidine bicarbonate in a mixture of a water-soluble solvent and polyphosphoric acid.

15 [0012] Suitably the total number of carbon atoms in R_1 to R_5 is less than eight.

[0013] The phenyl ring will suitably contain up to three substituents and preferably one or two substituents.

[0014] R_1 to R_5 are preferably independently selected from halogen and hydrogen. Particularly preferred substitutions are 4 or 2,3 or 2,4 or 3,4 mono- or di- halo (especially chloro).

20 [0015] Preferred compounds are:

- 4- Chlorobenzoyl cyanide amidinohydrazone,
- 2,3-Dichlorobenzoyl cyanide amidinohydrazone,
- 2,4- Dichlorobenzoyl cyanide amidinohydrazone,
- 3,4- Dichlorobenzoyl cyanide amidinohydrazone.

25 [0016] The preparation of the compounds of the formula (II) is analogous to that described in the literature, i.e. U.S. Patent No. 3,637,688; R.W.A. Rees et al, J. Med. Chem., 1972, 15, 859.

[0017] The amount of aminoguanidine bicarbonate is preferable more than one equivalent and more preferable 1.5 molar equivalents relative to compound of the formula (II).

30 [0018] The amount of polyphosphoric acid is preferably from 4 to 8 g. and more preferably 5.5 g to one gram of the compound of formula (II).

[0019] The reaction solvents are preferably water-soluble solvents such as acetonitrile, tetrahydrofuran, 1,2-dimethoxyethane (monoglyme), 2-methoxyethyl ether (diglyme) and the like and more preferably acetonitrile.

35 [0020] The amount of the reaction solvent is preferably from 2 to 8 ml, and more preferably 5 ml to one gram of the compound of the formula (II).

[0021] The reaction temperature may be chosen from ambient temperature to the reflux temperature of the solvent, and more preferably the temperature is between 50-70°C.

[0022] Since the progress of the reaction can be monitored by using high performance liquid chromatography, the reaction may be stopped after the disappearance of the starting material.

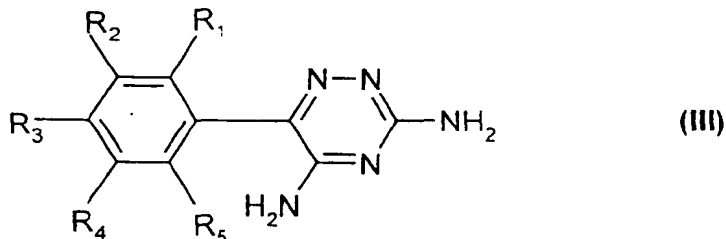
40 [0023] Although amidinohydrazones of the formula (I) exist as two isomers (E-and Z-forms), for the purpose of the present invention, these compounds may exist as either of the isomers or as a mixture thereof.

[0024] After completion of the reaction, water is added to the reaction mixture to obtain a clear homogeneous mixture. The mixture is dropped to water, and a colourless precipitate is collected by filtration to obtain a wet phosphate of the compound of the formula (I).

45 [0025] The wet phosphate of the compound of the formula (I) is added to an aqueous alkali solution, e.g., sodium hydroxide and the like, and the mixture is stirred at ambient temperature for one hour. A precipitate is then collected by filtration, washed with water and dried at 80°C to give a free base of the compound of the formula (I).

Triazines of the general formula (III):

[0026]



wherein R_1 to R_5 are as defined in the formula (I), can be obtained from the compound of the formula (I) by a procedure similar to that described in the literature, i.e. Beyer et al, *Chem. Ber.* 1960, **93**, 2209; J.A. Settepani and A.B. Borkovec, *J. Heterocycl. Chem.*, 1966, **3**, 188, U.S. Patent Nos. 3,637,688; R.W.A. Rees et al, *J. Med. Chem.*, 1972, **15**, 859; European Patent Nos. 21,121; 247,892.

[0027] According to the present invention, the compounds of the formula (I) can be obtained in high yield from the compound of the formula (II) by using polyphosphoric acid and simple procedures.

[0028] The present invention will be concretely illustrated by Exam, which show the method for preparation of the compound of formula (I).

[0029] While the invention will now be described in connection with certain preferred embodiments in the following examples so that aspects thereof may be more fully understood and appreciated, it is not intended to limit the invention to these particular embodiments. On the contrary, it is intended to cover all alternatives, modifications and equivalents as may be included within the scope of the invention as defined by the appended claims. Thus, the following examples which include preferred embodiments will serve to illustrate the practice of this invention, it being understood that the particulars shown are by way of example and for purposes of illustrative discussion of preferred embodiments of the present invention only and are presented in the cause of providing what is believed to be the most useful and readily understood description of formulation procedures as well as of the principles and conceptual aspects of the invention.

EXAMPLE 1:

Preparation of 2,3-Dichlorobenzoyl cyanide amidinohydrazone

[0030] a) Reactor (0.5 L) was charged under nitrogen with acetonitrile (200 ml) and polyphosphoric acid (220 g), and the mixture was stirred for 0.5 hour to obtain an emulsion. Solid 2,3-dichlorobenzoyl cyanide (41 g, 0.2 mole; assay 97.5%) was added to the emulsion. Aminoguanidine bicarbonate (41 g, 0.3 mole; assay 99.0%) was added to the mixture in 5 equal portions during 2 hours at 30-40°C. The reaction mixture was stirred at 50°C during 22 hours. Water (40 ml) was then added to the reaction mixture to obtain a clear homogeneous mixture. The mixture was dropped to water at slow stirring for 0.5 hour, and the precipitate was collected by filtration to give wet phosphate salt of 2,3-dichlorobenzoyl cyanide amidinohydrazone.

[0031] The wet phosphate salt was added at ambient temperature to 2.5% aqueous sodium hydroxide solution (500 ml), and the mixture was stirred for 0.5 hour. The precipitate was collected by filtration, washed thoroughly with water and dried at 80°C overnight to give 43.0 g (84% yield) of 2,3-dichlorobenzoyl cyanide amidinohydrazone base.

[0032] m.p. 221-222°C (uncorrected) (after dissolution in tetrahydrofuran, filtration of the solution and removal of the solvent to dryness *in vacuo*).

$^1\text{H NMR}$ (DMSO- d_6): δ (ppm) = 6.7 (4H, s, NH), 7.3-7.7 (3H, m, ArH).

$^{13}\text{C NMR}$ (DMSO- d_6): δ (ppm) = 113.88 and 114.53 ($\text{C}=\text{N}$, d, E- and Z-forms of the hydrazone), 128.24, 129.53 and 130.10 ($\text{C}-\text{H}$ from Ar), 163.64 [$-\text{NH}-\text{C}(\text{NH}_2)=\text{NH}$].

MS (FAB): m/z 256 (MH^+).

Analysis: Calcd. for $\text{C}_9\text{H}_7\text{Cl}_2\text{N}_5$: C, 42.21; H, 2.75; N, 27.35, found: C, 42.12; H, 2.77; N, 27.80.

[0033] b) The title compound was obtained in 67% yield after carrying out the reaction of 2,3-dichlorobenzoyl cyanide with aminoguanidine in a mixture of monoglyme and polyphosphoric acid.

EXAMPLE 2:

Preparation of 3,5-diamino-6-(2,3-dichlorophenyl)-1,2,4-triazine

[0034] a) The 2,3-dichlorobenzoyl cyanide amidinothydrazone base (43.0 g) obtained in Example 1(a) was heated under reflux in 1-propanol (560 ml) for two hours. The hot solution was filtered off, and the filtrate was cooled to 10°C. The crystals were collected by filtration, recrystallized from 1-propanol (490 ml) and dried at 100°C to give 3,5-diamino-6-(2,3-dichlorophenyl)-1,2,4-triazine (34.5 g).

[0035] The 1-propanol was evaporated to dryness *in vacuo* from the two filtrates, and the residue was recrystallized from 1-propanol to give 4.0 g of the title compound.

[0036] Total yield (from 2,3-dichlorobenzoyl cyanide): 38.5 g (75.2%); m.p. 216-218°C (uncorrected).

¹H NMR (DMSO-d₆): δ (ppm) = 6.16(2H, s, NH), 6.45 (2H, br s, NH), 7.2-7.7 (3H, m, ArH).

¹³C NMR (DMSO-d₆): δ (ppm) = 129.26, 131.35 and 131.51 (3 C-H from Ar), 155.114 [C-C(NH₂)=N-], 162.95 [=N-C(NH₂)=N-].

MS (FAB): *m/z* 256 (MH⁺).

[0037] b) Wet phosphate obtained by the procedure similar to that of Example 1(a) from 200 g (0.959 mole; assay: 95.91%) of 2,3-dichlorobenzoyl cyanide was added to 1-propanol (2.0 L), and the mixture was heated to 65°C at stirring. 45% Aqueous sodium hydroxide solution (185 ml) was then added to this mixture to produce pH 10, and the mixture was heated under reflux for two hours. The water phase was collected, and the organic phase was cooled to 10°C. The colourless crystals were collected by filtration, recrystallized from 1-propanol (2.0 L) (the hot solution was filtered) and dried at 100°C to give 160 g of the title compound.

[0038] An additional 20.0 g of the title compound was obtained from the filtrates. Total yield (from 2,3-dichlorobenzoyl cyanide): 180 g (73.3%).

[0039] c) The title compound was obtained in 60% total yield (from 2,3-dichlorobenzoyl cyanide) from 2,3-dichlorobenzoyl cyanide amidinothydrazone obtained in Example 1(b).

EXAMPLE 3:

Preparation of 2,4-dichlorobenzoyl cyanide amidinothydrazone

[0040] The title compound was prepared as in Example 1(a). Yield: 65%.

¹H NMR (DMSO-d₆): δ (ppm) = 6.70 (4H, s, NH), 7.4-7.8 (3H, m, ArH).

¹³C NMR (DMSO-d₆): δ (ppm) = 114.16 and 114.65 (C≡N, d, E- and Z-forms of the hydrazone), 128.06, 130.05 and 132.22 (3 C-H from Ar), 163.87 [-NH-C(NH₂)=NH].

MS (FAB): *m/z* 256 (MH⁺).

EXAMPLE 4:

Preparation of 3,5-Diamino-6-(2,4-dichlorophenyl)-1,2,4-triazine

[0041] The title compound was prepared as in Example 2(a) from 2,4-dichlorobenzoyl cyanide amidinothydrazone obtained in Example 3. The solvent for ring closure was a mixture of 1-propanol and dimethylsulfoxide (5:1). Total yield (from 2,4-dichlorobenzoyl cyanide): 55%; m.p. 220-222°C (uncorrected).

¹H NMR (DMSO-d₆): δ (ppm) = 6.40 (2H, s, NH), 6.66 (2H, br s, NH), 7.1-7.8 (3H, m, ArH).

¹³C NMR (DMSO-d₆): δ (ppm) = 127.60, 129.04 and 133.19 (3 C-H from Ar), 154.21 [C-C(NH₂)=N-], 161.98 [=N-C(NH₂)=N-].

MS (FAB): *m/z* 256 (MH⁺).

EXAMPLE 5:

Preparation of 3,4-Dichlorobenzoyl cyanide amidinothydrazone

[0042] The title compound was prepared as in Example 1(a). Yield: 86%.

¹H NMR (DMSO-d₆): δ (ppm) = 7.28 (4H, s, NH), 7.5-8.3 (3H, m, ArH). ¹³C NMR (DMSO-d₆): δ (ppm) = 112.44 and 117.20 (C≡N, d, E- and Z-forms of the hydrazone), 125.20, 126.06 and 131.95 (3 C-H from Ar), 160.99 [-NH-C(NH₂)=NH].

MS (FAB): *m/z* 256 (MH⁺).

EXAMPLE 6:**Preparation of 3,5-Diamino-6-(3,4-dichlorophenyl)-1,2,4-triazine**

[0043] The title compound was prepared as in Example 2(a) from 3,4-dichlorobenzoyl cyanide amidinohydrazone obtained in Example 5. The solvent for the ring closure was a mixture of 1-propanol and dimethyl sulfoxide (1:1). Total yield (from 3,4-dichlorobenzoyl cyanide): 76%; m.p. 222-224°C (uncorrected).

¹³C NMR (DMSO-d₆): δ (ppm) = 128.6, 130.24 and 130.83 (3 C-H from Ar), 155.02 [C-C(NH₂)=N-], 158.99 [=N-C(NH₂)=N-].

MS (FAB): *m/z* 256 (MH⁺).

EXAMPLE 7:**Preparation of 4-Chlorobenzoyl cyanide amidinohydrazone**

[0044] The title compound was prepared as in Example 1(a). Yield: 80%.

¹H NMR (DMSO-d₆): δ (ppm) = 6.70 (4H, s, NH), 7.2-8.0 (4H, m, ArH). ¹³C NMR (DMSO-d₆): δ (ppm) = 113.49 and 116.79 (C≡N, d, E- and Z-forms of the hydrazone), 125.95 and 128.44 (4 C-H from Ar), 163.90 [-NH-C(NH₂)=NH].

MS (FAB): *m/z* 222 (MH⁺).

EXAMPLE 8:**Preparation of 3,5-Diamino-6-(4-chlorophenyl)-1,2,4-triazine**

[0045] The title compound was prepared as in Example 2(a) from 4-chlorobenzoyl cyanide amidinohydrazone obtained in Example 7. The solvent for the ring closure was a mixture of 1-propanol and dimethyl sulfoxide (4:1). Total yield (from 4-chlorobenzoyl cyanide): 72%; m.p. 219-221°C (uncorrected).

¹H NMR (DMSO-d₆): δ (ppm) = 6.43 (2H, s, NH), 6.73 (2H, br s, NH), 7.3-7.7 (4H, m, ArH).

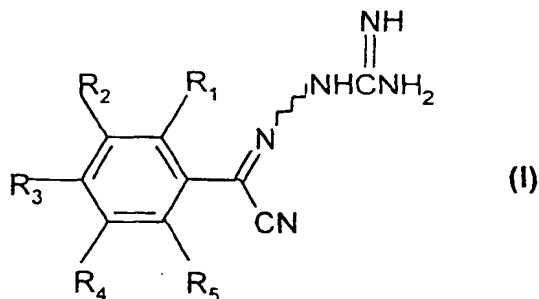
¹³C NMR (DMSO-d₆): δ (ppm) = 128.69 and 129.87 (4 C-H from Ar), 154.32 [C-C(NH₂)=N-], 161.55 [=N-C(NH₂)=N-].

MS (FAB): *m/z* 222 (MH⁺).

[0046] It will be evident to those skilled in the art that the invention is not limited to the details of the foregoing illustrative examples and that the present invention may be embodied in other specific forms without departing from the essential attributes thereof, and it is therefore desired that the present embodiments and examples be considered in all respects as illustrative and not restrictive, reference being made to the appended claims, rather than to the foregoing description, and all changes which come within the meaning and range of equivalency of the claims are intended to be embraced ther.

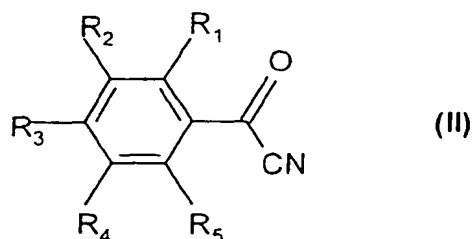
Claims

1. A process for preparing a compound of the general formula (I):



wherein R₁ to R₅ are independently selected from hydrogen, halogen, C₁₋₆ alkyl, C₂₋₆ alkenyl, C₂₋₆ alkynyl or C₁₋₆ alkoxy, all optionally substituted by one or more of halogen, hydroxy and aryl groups, as well as being independently selected from amino, mono- or disubstituted amino, alkenyloxy, acyl, acyloxy, cyano, nitro, aryl and alkylthio groups,

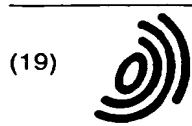
which process comprises reacting a compound of the general formula (II):



15 wherein R₁ to R₅ are as defined in formula (I), with aminoguanidine bicarbonate in a mixture of a water-soluble solvent and polyphosphoric acid.

- 20
2. A process according to claim 1 wherein the amount of aminoguanidine bicarbonate is more than 1 equivalent per 1 equivalent of the compound of formula (II).
 3. A process according to any one of claims 1 to 2 wherein the amount of polyphosphoric acid is from 4 to 8 g per one gram of the compound of formula (II).
 4. A process according to any one of claims 1 to 3 wherein said water-soluble solvent is acetonitrile.
 5. A process according to any one of claims 1 to 4 wherein the reaction of the compound of formula (II) with amino-
guanidine bicarbonate is carried out at a temperature of about 25-150°C.
 6. A process according to any one of claims 1 to 5 wherein the compound of formula (I) is 2,3-Dichlorobenzoyl cyanide
amidinohydrazone.
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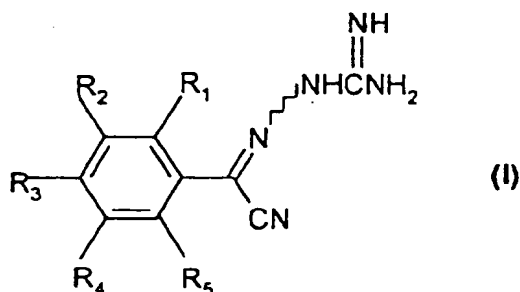
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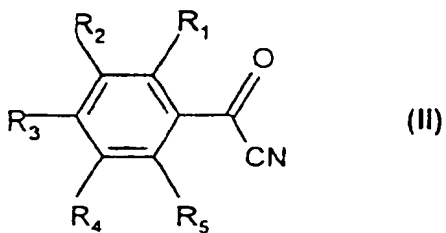
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(57) The present invention provides a process for preparing a compound of the general formula (I):



wherein R_1 to R_5 are independently selected from hydrogen, halogen, C_{1-6} alkyl, C_{2-6} alkenyl, C_{2-6} alkynyl or C_{1-6} alkoxy, all optionally substituted by one or more of halogen, hydroxy and aryl groups, as well as being independently selected from amino, mono- or disubstituted amino, alkenyloxy, acyl, acyloxy, cyano, nitro, aryl and alkylthio groups, which process comprises reacting a compound of the general formula (II):

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wherein R_1 to R_5 are as defined in formula (I), with aminoguanidine bicarbonate in a mixture of a water-soluble solvent and polyphosphoric acid.



European Patent
Office

EUROPEAN SEARCH REPORT

Application Number
EP 01 10 3660

DOCUMENTS CONSIDERED TO BE RELEVANT			
Category	Citation of document with indication, where appropriate, of relevant passages	Relevant to claim	CLASSIFICATION OF THE APPLICATION (Int.Cl.7)
D,A	EP 0 247 892 A (WELLCOME FOUND) 2 December 1987 (1987-12-02) examples 1 and 2 ---	1	C07C281/18
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